FDA 510(k) Premarket Notification

K001102 PARSET PRIMARY SET WITH CHECK VALVE, MODEL A10002E

CLEARED

Apr 20, 2000
15 days to decision

K001102 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k001102/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 5, 2000
Decision date	Apr 20, 2000
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Rd Medical Mfg., Inc.**
Location	Culebra, PR, US
Contact	CARLOS A RODRIGUEZ-GARCIA
510(k) history	4 submissions · 4 cleared · 1999-2001